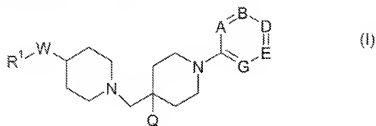


Amendments to the Claims:

This listing of claims replaces all prior versions and listings of claims in the application:

Listing of Claims:

1. (Original) A compound of formula (I):



wherein:

one of A, B, D, E and G is CXYCO₂R⁵, another is CH or N and the others are CR², CR³ and CR⁴;

Q is hydrogen or hydroxy;

W is CH₂, O, NH or N(C₁-₄ alkyl);

X is O or a bond;

Y is CR¹⁰R¹¹, CR¹⁰R¹¹CR¹²R¹³, CR¹⁰R¹¹CR¹²R¹³CR¹⁴R¹⁵;

R¹ is phenyl optionally substituted by halogen, cyano, C₁-₄ alkyl, C₁-₄ haloalkyl, C₁-₄ alkoxy or C₁-₄ haloalkoxy;

R², R³ and R⁴ are, independently, hydrogen, halogen, cyano, nitro, hydroxy, NR⁵R⁷, C₁-₆ alkyl (optionally substituted with halogen), C₁-₆ alkoxy (optionally substituted with halogen), S(O)ₚ(C₁-₆ alkyl), S(O)ₚCF₃ or S(O)₂NR⁸R⁹;

R⁵ is hydrogen, C₁-₆ alkyl or benzyl;

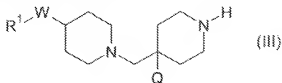
p and q are, independently, 0, 1 or 2;

R^6 , R^7 , R^8 and R^9 are, independently, hydrogen, C_{1-6} alkyl (optionally substituted by halogen, hydroxy or C_{3-6} cycloalkyl), $CH_2(C_{2-5}$ alkenyl), phenyl (itself optionally substituted by halogen, hydroxy, nitro, NH_2 , $NH(C_{1-4}$ alkyl), $N(C_{1-4}$ alkyl) $_2$ (and these alkyl groups may join to form a ring as described for R^6 and R^7 below), $S(O)_2(C_{1-4}$ alkyl), $S(O)_2NH_2$, $S(O)_2NH(C_{1-4}$ alkyl), $S(O)_2N(C_{1-4}$ alkyl) $_2$ (and these alkyl groups may join to form a ring as described for R^6 and R^7 below), cyano, C_{1-4} alkyl, C_{1-4} alkoxy, $C(O)NH_2$, $C(O)NH(C_{1-4}$ alkyl), $C(O)N(C_{1-4}$ alkyl) $_2$ (and these alkyl groups may join to form a ring as described for R^6 and R^7 below), CO_2H , $CO_2(C_{1-4}$ alkyl), $NHC(O)(C_{1-4}$ alkyl), $NHS(O)_2(C_{1-4}$ alkyl), $C(O)(C_{1-4}$ alkyl), CF_3 or OCF_3) or heterocyclyl (itself optionally substituted by halogen, hydroxy, nitro, NH_2 , $NH(C_{1-4}$ alkyl), $N(C_{1-4}$ alkyl) $_2$ (and these alkyl groups may join to form a ring as described for R^6 and R^7 below), $S(O)_2(C_{1-4}$ alkyl), $S(O)_2NH_2$, $S(O)_2NH(C_{1-4}$ alkyl), $S(O)_2N(C_{1-4}$ alkyl) $_2$ (and these alkyl groups may join to form a ring as described for R^6 and R^7 below), cyano, C_{1-4} alkyl, C_{1-4} alkoxy, $C(O)NH_2$, $C(O)NH(C_{1-4}$ alkyl), $C(O)N(C_{1-4}$ alkyl) $_2$ (and these alkyl groups may join to form a ring as described for R^6 and R^7 below), CO_2H , $CO_2(C_{1-4}$ alkyl), $NHC(O)(C_{1-4}$ alkyl), $NHS(O)_2(C_{1-4}$ alkyl), $C(O)(C_{1-4}$ alkyl), CF_3 or OCF_3); alternatively NR^6R^7 or NR^8R^9 may, independently, form a 4-7 membered heterocyclic ring, azetidine, pyrrolidine, piperidine, azepine, morpholine or piperazine, the latter optionally substituted by C_{1-4} alkyl on the distal nitrogen;

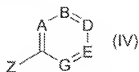
R^{10} , R^{11} , R^{12} , R^{13} , R^{14} and R^{15} are, independently, hydrogen or C_{1-4} alkyl; or R^{10} and R^{11} , and the carbon to which they are both attached, together form a C_{3-6} cycloalkyl ring, for C_{4-6} cycloalkyl rings said ring optionally having a ring carbon, but not the ring carbon to which R^{10} and R^{11} are both attached, replaced by O, $S(O)$ or $S(O)_2$; or an N-oxide thereof; or a pharmaceutically acceptable salt thereof.

2. (Original) A compound of formula (I) as claimed in claim 1 wherein W is O.

3. (Currently amended) A compound of formula (I) as claimed in claim 1-~~or~~2 wherein R^1 is phenyl optionally substituted with halogen, C_{1-4} alkyl or cyano.
4. (Currently amended) A compound of formula (I) as claimed in claim 1-~~2 or~~3 wherein R^2 , R^3 and R^4 are, independently, hydrogen, halogen, cyano, C_{1-4} alkyl, C_{1-4} alkoxy, CF_3 , OCF_3 , $S(O)_2(C_{1-4}$ alkyl) or $S(O)_2NH_2$.
5. (Currently amended) A compound of formula (I) as claimed in ~~any one of the preceding claims~~claim 1 wherein Q is hydrogen.
6. (Currently amended) A compound of formula (I) as claimed in ~~any one of the preceding claims~~claim 1 wherein one of A, B, D, E and G is $CXYCO_2R^5$ and the others are all CH.
7. (Currently amended) A compound of formula (I) as claimed in ~~any one of the preceding claims~~claim 1 wherein XY is CH_2 , CH_2CH_2 , OCH_2 , $OC(CH_3)_2$ or $OCHCH_3$.
8. (Currently amended) A compound of formula (I) as claimed in ~~any one of the preceding claims~~claim 1 wherein R^5 is hydrogen or C_{1-6} alkyl.
9. (Original) A process for preparing a compound of formula (I) as claimed in claim 1, the process comprising:
 - a. when R^5 is alkyl or benzyl, esterifying a compound of formula (I) where R^5 is H;
 - b. when R^5 is H, hydrolyzing a compound of formula (I) wherein one of A, B, D, E, or G is $CXYCN$;
 - c. reacting a compound of formula (III)

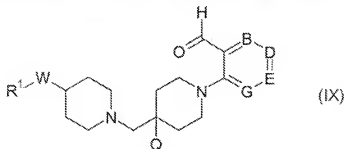


with a compound of formula (IV)



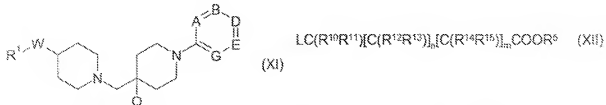
wherein Z is Br, I; in the presence of copper iodide, proline and a base in a suitable solvent at a suitably elevated temperature;

- d. reacting a compound of formula (III) with a compound of formula (IV), wherein Z is Br or I, in the presence of a palladium salt, a phosphine and a base, in a suitable solvent at a suitably elevated temperature;
- e. when A is $CXYCO_2R^5$, reacting a compound of formula (IX):



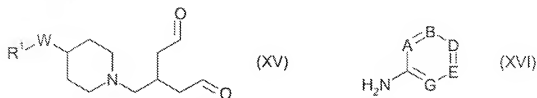
with methyl methylthiomethyl sulfoxide or ethyl ethylthiomethyl sulfoxide in the presence of a base, in a suitable solvent, at a suitable temperature, and treating the product resulting therefrom with HCl in R^5OH ;

- f. when XY is $OCR^{10}R^{11}$, $OCR^{10}R^{11}CR^{12}R^{13}$ or $OCR^{10}R^{11}CR^{12}R^{13}CR^{14}R^{15}$, reacting a compound of formula (XI), wherein one of A, B, D, E, or G represents $C(O)H$, with a compound of formula (XII), wherein L is halogen or a sulfonate ester, and n and m are, independently, 0 or 1,



in the presence of a base, in a suitable solvent at ambient temperature;

- g. when Q is H, reacting a compound of formula (XV) with a compound of formula (XVI)



in the presence of a suitable reducing agent and acetic acid, in a suitable solvent.

10. (Original) A pharmaceutical composition which comprises a compound of the formula (I), or a pharmaceutically acceptable salt thereof as claimed in claim 1, and a pharmaceutically acceptable adjuvant, diluent or carrier.
- 11-12. (Cancelled)
13. (Original) A method of treating a chemokine mediated disease state in a mammal suffering from, or at risk of, said disease, which comprises administering to a mammal in need of such treatment a therapeutically effective amount of a compound of formula (I), or a pharmaceutically acceptable salt thereof as claimed in claim 1.